

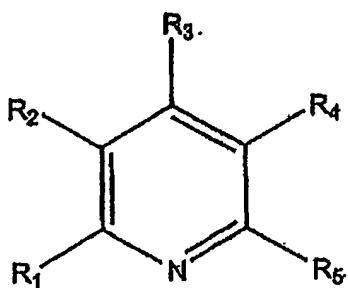
**Amendments to the Claims:**

This listing of claims will replace all prior versions and listing of claims in the application.

Claims 4-10, 24-25 and 28-44 are amended.

**Listing of Claims:**

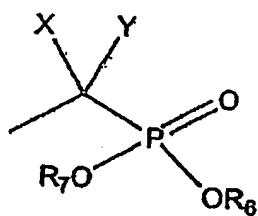
1. (Original) A compound of general formula:



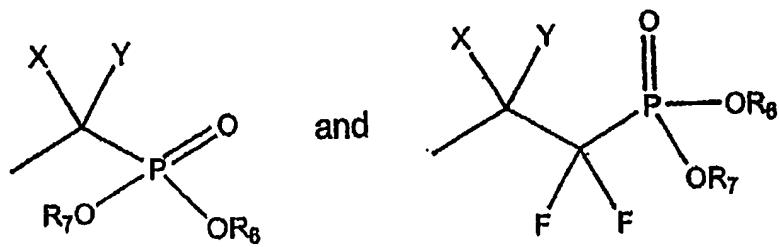
Wherein:

R<sub>1</sub> is selected from H and CH<sub>3</sub>, and R<sub>2</sub> is selected from H and OH, or R<sub>1</sub> and R<sub>2</sub> together form an optionally substituted phenyl ring which is fused to the pyridine ring; and

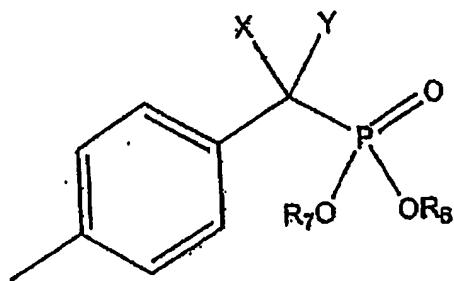
R<sub>3</sub> is selected from H, CH<sub>3</sub>, CH<sub>2</sub>OH and



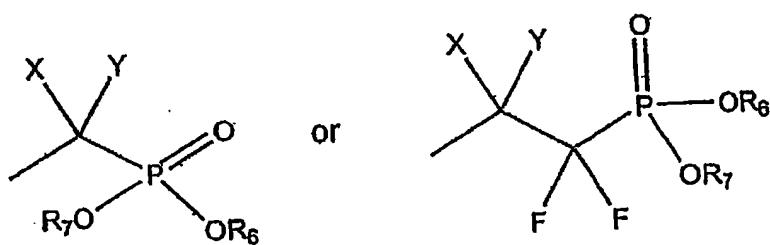
R<sub>4</sub> is selected from H, CH<sub>3</sub>, CH<sub>2</sub>OH,



$R_5$  is selected from H, phenyl, halogen-substituted phenyl and

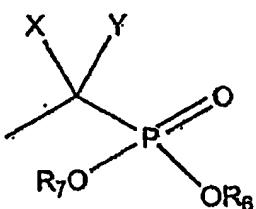


Wherein  $R_6$  and  $R_7$  are each independently selected from H,  $Na^+$ ,  $K^+$ , alkyl and optionally substituted aryl, and X and Y are each independently selected from H, OH and F, or at least one of X and Y is an heteroatom and together with  $R_3$  forms a bridge with the proviso that  $R_4$  is

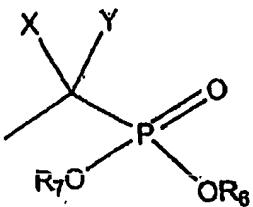


and N-oxides thereof, and biologically acceptable salts thereof.

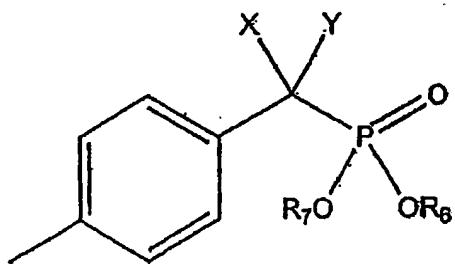
2. (Original) The compound according to claim 1, wherein said halogen-substituted phenyl is a fluoro-substituted phenyl.
3. (Original) The compound according to claim 1, wherein said halogen-substituted phenyl is p-C<sub>6</sub>H<sub>4</sub>F.
4. (Currently Amended) The compound according to ~~any one of claims 1 to 3~~ claim 1, wherein said heteroatom is selected from O and S.
5. (Currently Amended) The compound according to ~~any one of claims 1 to 3~~ claim 1, wherein said heteroatom is O.
6. (Currently Amended) The compound according to ~~any one of claims 1 to 5~~ claim 1, wherein said bridge is selected from -CH<sub>2</sub>-, -CH<sub>2</sub>CH<sub>2</sub>- and -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-.
7. (Currently Amended) The compound according to ~~any one of claims 1 to 5~~ claim 1, wherein said bridge is a methylene bridge.
8. (Currently Amended) The compound according to ~~any one of claims 1 to 7~~ claim 1, wherein said alkyl is a C<sub>1</sub> to C<sub>6</sub> straight or branched alkyl.
9. (Currently Amended) The compound according to ~~any one of claims 1 to 7~~ claim 1, wherein said alkyl is t-butyl.
10. (Currently Amended) The compound according to ~~any one of claims 1 to 9~~ claim 1, wherein said aryl is phenyl or naphthyl.
11. (Original) The compound according to claim 1, wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>5</sub> are all H and R<sub>3</sub> is



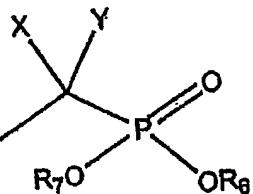
12. (Original) The compound according to claim 1, wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>5</sub> are all H and R<sub>4</sub> is



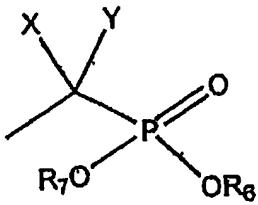
13. (Original) The compound according to claim 1, wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> are all H and R<sub>5</sub> is



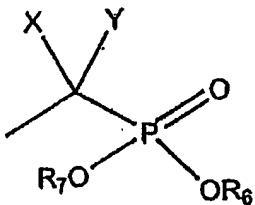
14. (Original) The compound according to claim 1 wherein R<sub>1</sub> and R<sub>2</sub> together form an optionally substituted phenyl ring which is fused to the pyridine ring; R<sub>3</sub> and R<sub>5</sub> are both H; and R<sub>4</sub> is



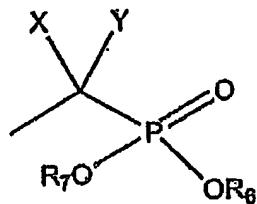
15. (Original) The compound according to claim 1, wherein R<sub>1</sub> and R<sub>3</sub> are both CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>5</sub> is H; and R<sub>4</sub> is



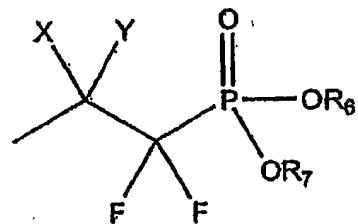
16. (Original) The compound according to claim 1, wherein R<sub>1</sub> and R<sub>4</sub> are both CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>5</sub> is H; and R<sub>3</sub> is



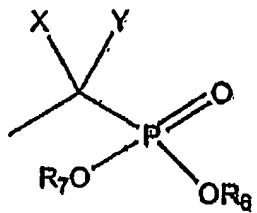
17. (Original) The compound according to claim 1, wherein R<sub>1</sub> is CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>3</sub> is CH<sub>2</sub>OH; R<sub>5</sub> is H; and R<sub>4</sub> is



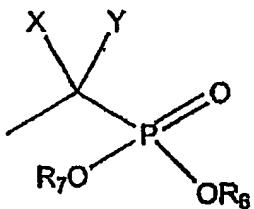
18. (Original) The compound according to claim 1, wherein R<sub>1</sub> is CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>3</sub> is CH<sub>2</sub>OH; R<sub>5</sub> is H; and R<sub>4</sub> is



19. (Original) The compound according to claim 1, wherein R<sub>1</sub> is CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>5</sub> is CH<sub>2</sub>OH; R<sub>5</sub> is C<sub>6</sub>H<sub>5</sub>; and R<sub>4</sub> is



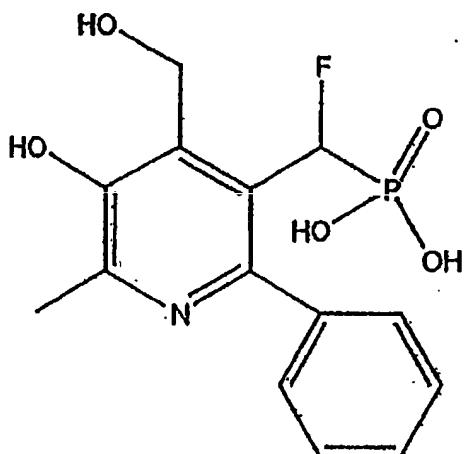
20. (Original) The compound of claim 1, wherein R<sub>1</sub> is CH<sub>3</sub>; R<sub>2</sub> is OH; R<sub>3</sub> is CH<sub>2</sub>OH; R<sub>5</sub> is p-C<sub>6</sub>H<sub>4</sub>F; and R<sub>4</sub> is



21. (Original) A compound according to claim 1, wherein R<sub>5</sub> is P-C<sub>6</sub>H<sub>4</sub>F.

22. (Original) A compound according to claim 1 selected from: [Hydroxy-(5-hydroxy-4-hydroxymethyl-6-methyl-2-phenyl-pyridin-3-yl)-methyl]-phosphonic acid; {[2-(4Fluoro-phenyl)-5-hydroxy-4-hydroxymethyl-6-methyl-pyridin-3-yl]-hydroxymethyl}phosphonic acid; [Hydroxy-(4-pyridin-2-yl-phenyl)-methyl]-phosphonic acid; [Fluoro(4-pyridin-2-yl-phenyl)-methyl]-phosphonic acid; (Hydroxy-quinolin-3-yl-methyl)phosphonic acid; (Fluoro-quinolin-3-yl-methyl)-phosphonic acid; [Hydroxy-(5hydroxy-4, 6-dimethyl-pyridin-3-yl)-methyl]-phosphonic acid; (Hydroxy-pyridin-4-yl-methyl)-phosphonic acid; (Hydroxy-pyridin-3-yl-methyl)-phosphonic acid; (3,7Dihydroxy-6-methyl-1,3-dihydro-furo[3,4-c]pyridin-3-yl)-phosphonic acid; [(3,7Dihydroxy-6-methyl-1,3-dihyrdio-furo[3,4-c]pyrid i n-3-yl )-difluoromethyl]-phosphonic acid; and nicotinyl phosphonates thereof, N-oxides thereof, phosphonate esters thereof and biologically acceptable salts thereof.

23. (Original) A compound according to claim 1 comprising:



24. (Currently Amended) A pharmaceutical composition comprising a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 23~~ claim 1 and a pharmaceutically acceptable carrier.

25. (Currently Amended) The compound according to ~~anyone of claims 1 to 24~~ claim 1, wherein at least one polar group is blocked by a lipophilic moiety capable of being enzymatically cleaved off after absorption into the circulatory system.

26. (Original) The compound according to claim 25, wherein said lipophilic moiety is an ester.

27. (Original) The compound according to claim 25, wherein said lipophilic moiety is a phosphonate ester.

28. (Currently Amended) A method of treating hypertension in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

29. (Currently Amended) A method of treating myocardial infarction in a mammal

comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

30. (Currently Amended) A method of treating ischemia reperfusion injury in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

31. (Currently Amended) A method of treating myocardial ischemia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

32. (Currently Amended) A method of treating congestive heart failure in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

33. (Currently Amended) A method of treating arrhythmia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

34. (Currently Amended) A method of reducing blood clots in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

35. (Currently Amended) A method of treating hypertrophy in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

36. (Currently Amended) A method of treating a disease that arises from thrombotic and

prothrombotic states in which the coagulation cascade is activated in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

37. (Currently Amended) A method of treating diabetes mellitus in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

38. (Currently Amended) A method of treating insulin resistance in a mammal comprising concurrently administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

39. (Currently Amended) A method of treating hyperinsulinemia in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ in a unit dosage form.

40. (Currently Amended) A method of treating diabetes-induced hypertension in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

41. (Currently Amended) A method of treating diabetes-related damage to blood vessels, eyes, kidneys, nerves, autonomic nervous system, skin, connective tissue, or immune system in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

42. (Currently Amended) A method of treating obesity in a mammal comprising administering to the mammal a therapeutically effective amount of a compound according to ~~anyone of claims 1 to 27~~ claim 1 in a unit dosage form.

43. (Currently Amended) A compound according to ~~anyone of claims 1 to 27~~ claim 1 which is a nicotinic acid derivative.

44. (Currently Amended) A kit comprising the composition of ~~anyone of claims 1 to 27~~ claim 1 and instructions for its use in the treatment of a cardiovascular disease, a disease that arises from a thrombotic or prothrombotic state in which the coagulation cascade is activated, diabetis, or related diseases.